

The Management of osteomyelitis by stimulant.

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Abstract

The complex and heterogeneous nature of chronic osteomyelitis, necessitates a multi-disciplinary approach, involving experts in the field of orthopaedic tumour, infection and limb reconstruction surgery, plastic surgery, microbiology and nursing. Numerous surgical techniques and adjuvant therapies have been developed during the past three decades in order to deal with the wide spectrum of pathology that falls under the heading of chronic osteomyelitis. Despite these developments, the outcome of current treatment protocols remains unsatisfactory, with failure of therapy. Cases with osteomyelitis of long bone in adolescents which were previously managed by multiple surgeries, presented with recurrence, discharging sinus, pus was first sent for culture and found to be *Staphylococcus aureus* sensitive to Vancomycin. Sequestrectomy and saucerisation and thorough debridement of the canal was performed following which Stimulan pellets prepared with vancomycin were packed in the canal, wound was closed. Serial monitoring was done once in 3 weeks after about 2 months xray showed a good response. Since biodegradable antibiotic drug delivery systems are effective against osteomyelitis, it stands to reason that they might also be effective in prophylaxis.

Keywords: Orthopaedic tumour, Seuestrectomy.

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Introduction

The use of local antibiotics to treat osteomyelitis effectively necessitates a thorough grasp of the disease's features and pathophysiology. *Staphylococcus* species are the most prevalent pathogens responsible for osteomyelitis in humans, followed by Enterobacteriaceae and *Pseudomonas* species. More than 90% of the time, *Staphylococcus epidermidis* causes osteomyelitis associated with an implant. 1 Diabetic foot infections are typically polymicrobial infections including a mixture of gram-positive and gram-negative bacteria, as well as aerobic and anaerobic bacteria. Chronic osteomyelitis has a complex pathogenesis that begins with bacterial spread. Hematogenous seeding, direct inoculation, or airborne contamination can all cause bacteria to enter the bone. Bacteria have a variety of methods for facilitating cell-cell and cell-implant adhesion once they come into contact with bone and/or implant. Bacteria become sessile, slow down their metabolism, and form a biofilm to defend themselves against antimicrobials, opsonization, and phagocytosis. Adhesins in *Staphylococcus aureus*, for example, bind fibronectin. Fibronectin, a connective tissue glycoprotein, has been demonstrated to mediate and enhance bacterial-foreign-body interactions² Antibiotics work on multiple levels to prevent bacterial illness. Clindamycin, for example, inhibits glycocalyx formation, allowing phagocytic cells to function more effectively and decreasing infection by glycocalyx-forming organisms. 3 Clindamycin also inhibits bacterial protein synthesis by preferentially binding to the 50S component of bacterial ribosomes.

Antibiotic concentrations required to penetrate and kill bacteria encased in biofilm, on the other hand, are 10 to 100 times higher than typical bactericidal concentrations, rendering systemic therapy in such instances dangerous and useless. Sequestra, avascular areas of necrotic cortical bone that are difficult to heal, are caused by a prolonged bacterial infection of the bone that inhibits cortical blood supply. The sequestra is frequently

surrounded by involucrum, which is new bone formed in response to the sequestra. 5 The presence of the sequestra warrants surgical intervention in chronic osteomyelitis. Intravenous antibiotics are frequently effective in treating acute illnesses. A variety of conditions might cause a bone infection to become chronic.

Continued infection is aided by the presence of foreign substances and necrotic bone. Furthermore, pathogen-specific traits such as bacteria's capacity to remain intracellular, build a protective layer, and maintain a slow metabolic rate all impede the host's ability to remove infection. Obesity and smoking are two issues that affect patients. Chronic osteomyelitis is characterised by recurrent or intermittent disease, as well as the occurrence of late infection reactivation.

Latent periods can endure for decades, and some have been known to last up to 80 years. 6 The radiographic symptoms of osteomyelitis can be faint, thus they must be compared to the clinical picture. On routine radiographs, loss of trabecular marks, elevation of the periosteum, cortical scalloping, medullary involvement, and localised osteopenia are all signs of osteomyelitis. A sequestrum with surrounding involucrum may be visible on computed tomography (CT) images.

A subperiosteal fluid collection as well as a Brodie abscess, a subacute or chronic metaphyseal abscess of a bone that occurs as a pus-filled cavity surrounded by a wall of dense fibrous tissue and is most commonly found in the metaphyses of the long bones, can be seen on magnetic resonance imaging (MRI)

With the use of flaps and vascularized bone grafts, the surgical management of chronic osteomyelitis has altered dramatically in the last 25 years. Surgical intervention is nearly always required due to a lack of blood supply to the affected areas and a decreased ability of the host immune system to clear infections. In chronic osteomyelitis, thorough bone debridement is critical, and it is frequently the most important aspect in eradicating infection. have a negative impact on the treatment At baseline and at follow-up, pre- and post-operative data on the osseointegrated prostheses were compared for each patient. Between 1999 and 2007, 51 TFA patients were enrolled in the trial, with each patient being followed for five years.

References

1. Ciampolini J, Harding K G. Pathophysiology of chronic bacterial osteomyelitis. Why do antibiotics fail so often? *Postgrad Med J.* 2000;76:479–483
2. Fischer B, Vaudaux P, Magnin M, et al. Novel animal model for studying the molecular mechanisms of bacterial adhesion to bone-implanted metallic devices: role of fibronectin in *Staphylococcus aureus* adhesion. *J Orthop Res.* 1996;14:914–920.
3. Adams K, Couch L, Cierny G, Calhoun J, Mader J T. In vitro and in vivo evaluation of antibiotic diffusion from antibiotic-impregnated polymethylmethacrylate beads. *Clin Orthop Relat Res.* 1992;278:244–252.
4. Nelson C L. The current status of material used for depot delivery of drugs. *Clin Orthop Relat Res.* 2004;427:72–78.

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